

# Effects of rainfall on *Culex* mosquito population dynamics

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## Abstract

The dynamics of a mosquito population depends heavily on climatic variables such as temperature and precipitation. Since climate change models predict that global warming will impact on the frequency and intensity of rainfall, it is important to understand how these variables affect the mosquito populations. We present a model of the dynamics of a *Culex quinquefasciatus* mosquito population that incorporates the effect of rainfall and use it to study the influence of the number of rainy days and the mean monthly precipitation on the maximum yearly abundance of mosquitoes  $M_{max}$ . Additionally, using a fracturing process, we investigate the influence of the variability in daily rainfall on  $M_{max}$ . We find that, given a constant value of monthly precipitation, there is an optimum number of rainy days for which  $M_{max}$  is a maximum. On the other hand, we show that increasing daily rainfall variability reduces the dependence of  $M_{max}$  on the number of rainy days, leading also to a higher abundance of mosquitoes for the case of low mean monthly precipitation. Finally, we explore the effect of the rainfall in the months preceding the wettest season, and we obtain that a regimen with high precipitations throughout the year and a higher variability tends to advance slightly the time at which the peak mosquito abundance occurs, but could significantly change the total mosquito abundance in a year.

**Keywords:** *Culex*, rainfall, arbovirus, mosquito abundance, mathematical modeling

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## 1. Introduction

Mosquito-transmitted flaviviruses are an increasing health threat. In particular, members of *Culex* species (Diptera: Culicidae) such as *Cx. quinquefasciatus* and *Cx. interfor*

are responsible for transmitting the West Nile and St. Louis encephalitis (SLEV) viruses to humans and domestic animals (Beltrán et al. (2015); Diaz et al. (2008, 2016); Gubler (2002); Gubler et al. (2007); Lumsden (1958)). For instance, the SLEV is endemic in Argentina, where the principal vector is postulated to be *Cx. quinquefasciatus* (Diaz et al. (2013)). In the last decades, mosquito-borne diseases have emerged and re-emerged as a result of multiple factors such as increasing urbanization, international travel, and climate change (Harrigan et al. (2014); Kilpatrick (2011)). The development of mathematical models is essential to quantify the effect of each of these factors on the dynamics of the mosquito population, and to determine the most effective strategies to control the epidemic outbreaks transmitted by mosquito vectors (Anderson et al. (1992); Ewing et al. (2016); Lord and Day (2001a,b); Marini et al. (2016)).

Multiple studies have shown that the life cycle of the *Cx. quinquefasciatus* is closely related to temperature (Almirón and Brewer (1996); Ciota et al. (2014); Gunay et al. (2011); Loetti et al. (2011); Strickman (1988)). Strickman (1988) demonstrated that its reproductive activity increases with temperature, and Almirón and Brewer (1996) showed that this species can only live in environments with a temperature above 10°C. Other studies have found that temperature has a strong influence on the development and survival of both adult and immature mosquitoes (Ciota et al. (2014); Gunay et al. (2011); Loetti et al. (2011)). In turn, it was observed that *Cx. quinquefasciatus* does not enter diapause, but it may undergo quiescence or remain gonoactive in protected (indoor or underground) habitats (Almirón and Brewer (1996); Nelms et al. (2013)). Urbanization therefore helps *quinquefasciatus* populations survive the mild winters of temperate regions.

Similarly, rainfall is an important climatological variable to predict the abundance of *Culex* mosquitoes, since its copiousness and distribution determine the production and size of mosquito breeding sites. Reisen et al. (2008) studied the changes in the *Cx. tarsalis* population in California and found that, in most regions, it is positively correlated with an increase in total precipitation. However, these authors also found that in some places of the driest region of California, the correlation between these two variables was negative. On the other hand, Olson et al. (1983) showed that a very large rainfall is not always accompanied by proportionately large increases in the abundance of

*Cx. tritaeniorhynchus* and *Cx. gelidus*. In consequence, these results indicate that there exists a nonlinear relationship between rainfall and *Culex* abundance, which should be modeled in order to predict mosquito abundance. Additionally, since the climatological projections suggest that global warming will alter the frequency and intensity of rainfall, it is crucial to understand how different rainfall patterns will affect mosquito populations.

In this paper we develop a dynamic model of the *Cx. quinquefasciatus* population, adapting a fracturing procedure (Finley and Kilkki (2014)) to describe the rainfall distribution. We use a system of compartmental ordinary differential equations that describe the immature and adult mosquito populations, in which we introduce the influence of temperature and rainfall on the reproduction rate. To study the influence of different rainfall patterns, we use a synthetic time series of rainfall based on the amount of rainfall per month and the monthly number of rainy days. We find that, for a given constant value of monthly precipitation, there is an optimum number of rainy days for which the maximum  $M_{max}$  in the mosquito population is highest.

On the other hand, we also study the variability of daily rainfall intensity through a fracturing process (Finley and Kilkki (2014)), which allows us to study homogeneous and heterogeneous rainfall regimes, including those characterized by heavy rain events. We show that increasing daily rainfall variability reduces the dependence of  $M_{max}$  on the number of rainy days, leading also to a higher abundance of mosquitoes for the case of low mean monthly precipitation.

Finally, we explore the effect of different winter precipitation regimes on the mosquito abundance in the summer season, obtaining that a higher variability tends to advance slightly the peak time of mosquito abundance. Interestingly, we predict that the accumulated abundance of mosquitoes will decrease in a regime with high variability in the rainfall intensity.

The boundary of the *Cx. quinquefasciatus* habitat in South America runs across central Argentina. This region is thus expected to exhibit intense changes in mosquito populations due to the undergoing climatic change; this is likely to have a strong impact on flavivirus prevalence. For this reason, we use the climatic data for the city of Córdoba to calibrate our model in the period 2008-2009.

The paper is organized as follows: in Sec. 2.1 we present the model of the dynamics of mosquito population and in Secs. 2.2 and 2.3 we explain the methods for generating two different synthetic time series. Then in Sec. 3 we show our results and finally we present our conclusions in Sec. 4.

## 2. Methods

### 2.1. The model of mosquito abundance

In this section, we construct a compartmental, ordinary differential equation model for the mosquito abundance. We consider that the total vector population is stage-structured with an immature class consisting of all aquatic stages, and a mature or adult class. We assume that these population groups are restricted only to female mosquitoes as the reproductive sex.

In our model, the total birth rate, *i.e.* the total number of new immature female mosquitoes per unit of time, is proportional to the number of adult mosquito females and to  $\beta_L \lambda(t) \theta(t)$ , where  $\beta_L$  corresponds to the reproduction rate in optimal conditions of temperature and water availability, and  $\lambda(t)$  and  $\theta(t)$  are normalized factors describing the influence of rainfall and temperature on the total birth rate, respectively. In Sec. 2.1.1 we will explain how we construct these factors. In addition, we assume that the total birth rate is also regulated by a carrying capacity effect that depends on the occupation of the available immature habitats. Therefore we propose that the immature population growth is logistic-like with a carrying capacity  $K_L$ . Additionally, immature individuals either go to the mature class with rate  $m_L$  or die at a rate  $\mu_L$ . We stress that the ratio  $1/m_L$  gives the average development time from immature mosquito to adult. Finally, adult mosquitoes die with a mortality rate  $\mu_M$ . For simplicity, we assume that  $m_L$ ,  $\mu_L$  and  $\mu_M$  do not depend on temperature or rainfall. With these definitions, we propose the following dynamic mass-balance equations for the abundance of immature mosquitoes,  $L(t)$ , and adult mosquitoes,  $M(t)$ ,

$$L(t + \Delta t) = L(t) + \Delta t \left[ \beta_L \theta(t) \lambda(t) M(t) \left( 1 - \frac{L(t)}{K_L} \right) - m_L L(t) - \mu_L L(t) \right], \quad (1)$$

and

$$M(t + \Delta t) = M(t) + \Delta t [m_L L(t) - \mu_M M(t)], \quad (2)$$

where  $\Delta t$  is the time step size. Here we use  $\Delta t = 0.1$  [days]. Equation (1) can be easily derived by assuming that the fraction of the carrying capacity corresponding to female immature mosquitoes is the same as the fraction of females in the immature population. Table 1 summarizes the different state variables and parameters used in this paper.

Table 1: The variables and parameters for Eqs. (1)-(9).

Quantity	Definition	Value	Refs.
$L$	number of immature female mosquitoes	—	fitted (see Appendix A)
$M$	number of adult female mosquitoes	—	
$\beta_L$	birth rate of immature female mosquito per female adult mosquito in optimal conditions of temperature and water availability ( $\text{days}^{-1}$ )	13.5	
$\theta$	effect of the temperature on the birth rate of mosquitoes	—	
$\lambda$	effect of the water availability on the birth rate of mosquitoes	—	
$m_L$	rate at which immature mosquitoes develop into adults ( $\text{days}^{-1}$ )	0.098	
$\mu_L$	immature mortality rate ( $\text{days}^{-1}$ )	0.03	Loetti et al. (2011)
$\mu_M$	mosquito mortality rate ( $\text{days}^{-1}$ )	0.078	David et al. (2012)
$K_L$	carrying capacity of the immature female population	16.2	fitted (see Appendix A)
$H_{max}$	maximum daily amount of accumulated rainwater [mm]	9.86	fitted (see Appendix A)
$H_{min}$	minimum daily amount of accumulated rainwater [mm]	0.067	fitted (see Appendix A)
$H$	accumulated amount of rainwater [mm]	—	
$R$	daily rainfall [mm]	—	
$E$	daily evapotranspiration [mm]	—	
$T$	average daily temperature [ $^{\circ}\text{C}$ ]	—	
$Hum$	daily relative humidity	—	
$P_{max}$	total rainfall in the wettest month [mm]	—	
$P_{min}$	total rainfall in the driest month [mm]	—	
$D_{max}$	total number of rainy days in the wettest month [days]	—	
$D_{min}$	total number of rainy days in the driest month [days]	—	

### 2.1.1. Effect of temperature and rainfall on the total birth rate

We add the effect of the temperature through a temperature factor  $\theta(t)$ . Several studies have shown that *Cx. quinquefasciatus* can breed only at temperatures above 10°C (Almirón and Brewer (1996); Ribeiro et al. (2004)) and that the number of egg rafts collected per day is closely correlated with temperature (Strickman (1988)). Therefore we propose that the factor  $\theta(t)$  is a piecewise linear function,

$$\theta(t) = \begin{cases} \frac{T(t)-T_A}{T_B-T_A} & \text{if } T_A \leq T(t) \leq T_B \\ 1 & \text{if } T(t) > T_B \\ 0 & \text{if } T(t) < T_A, \end{cases} \quad (3)$$

where  $T(t)$  is the average daily temperature and  $T_A = 10^\circ\text{C}$  corresponds to a minimum temperature below which the net birth rate vanishes. The choice of the function  $\theta(t)$  is not unique: for instance, a power law may be used instead (Ewing et al. (2016)). Here we assume that the positive effect of the temperature on the birth rate saturates at  $T_B$ , since it was observed by Oda et al. (1980) that the number of egg rafts per female does not significantly change between 21°C and 30°C.

On the other hand, mosquito reproduction is also triggered by rainfalls since these increase the number of breeding sites, such as temporary ground pools. In order to introduce the effect of rainfall on the mosquito birth rate, we compute the accumulated amount of rainwater  $H$ , whose variation is given by the total daily rainfall  $R(t)$  minus the evapotranspiration  $E(t)$  (Gong et al. (2011)),

$$H(t+1) = H(t) + [R(t) - E(t)]. \quad (4)$$

The function  $H(t)$  is a convenient measure of the quantity of water available for breeding sites. It should represent the average level of puddles, ponds, drains, small streams, and underground sources such as waste water channels in urban environments. As we will explain below, equation (4) is applied on a daily time scale. The term of evapotranspiration is estimated using the Ivanov model (Romanenko (1961); Valipour (2014)), which is based on the mean temperature and the relative humidity  $[Hum(t)]$ , and it is given by

$$E(t) = 6.10^{-5}(25 + T(t))^2(100 - Hum(t)). \quad (5)$$

Note that the evapotranspiration  $E(t)$  is a monotonically increasing function of temperature and a decreasing function of humidity. In particular,  $E(t)$  vanishes for  $Hum = 100\%$ . In addition, we assume that the level of accumulated water  $H$  varies only between a minimum ( $H_{min}$ ) and a maximum ( $H_{max}$ ) boundary level, *i.e.*,

$$H(t+1) = \begin{cases} H_{min} & \text{if } H(t) + [R(t) - E(t)] \leq H_{min} \\ H_{max} & \text{if } H(t) + [R(t) - E(t)] \geq H_{max} \\ H(t) + [R(t) - E(t)] & \text{otherwise.} \end{cases} \quad (6)$$

Here,  $H_{min}$  represents a minimum amount of water that is always available for mosquito breeding, for instance in permanent streams or in the drainage system, while  $H_{max}$  is a level of water above which the breeding sites overflow (Karl et al. (2014)).

Finally, the factor  $\lambda(t)$  that takes into account the effect of rainfall on the birth rate (see Eq. (1)), is the normalization of  $H(t)$ :

$$\lambda(t) = \frac{H(t)}{H_{max}}. \quad (7)$$

Note that the minimum value of  $\lambda(t)$  is  $H_{min}/H_{max}$ .

Although we take the integration time step to be  $\Delta t = 0.1$  days, the data of temperature, rainfall and humidity are available only on a daily time scale. Therefore, for all the integration time steps within a day “ $d$ ” (*i.e.*  $d \leq t < d+1$ ), we set  $\lambda(t)$  and  $\theta(t)$  to have the values computed through Eqs. (6) and (5) using the corresponding meteorological data for day “ $d$ ”.

## 2.2. Model of synthetic rainfall using the monthly number of rainy days and the monthly precipitation

The amount of average rainfall  $P$  and the number of days  $D$  with rain per month are two parameters commonly used to characterize the long-term precipitation trend (Madsen et al. (2009); Owusu and Waylen (2013); Zhai et al. (2005)). In this section, we show how they can be used to construct a synthetic rainfall time series.

The average monthly rainfall is assumed to follow a sinusoidal function,

$$P(m) = \frac{P_{max} - P_{min}}{2} \cos\left(\frac{2\pi}{12}(m - m_0)\right) + \frac{P_{max} + P_{min}}{2}, \quad (8)$$



where  $m = 1, \dots, 12$  represents the month (with  $m = 1$  for January and  $m = 12$  for December),  $m_0$  corresponds to the month of maximum rainfall,  $P_{max}$  is the total precipitation of the wettest month  $m_0$ , and  $P_{min}$  is the total precipitation of the driest month, corresponding to  $m = m_0 + 6$ . Note that for a higher value of either  $P_{max}$  or  $P_{min}$  there is an increase in the annual amount of precipitation but, while a higher  $P_{max}$  enhances the precipitation difference between the rainy and dry seasons, a higher  $P_{min}$  reduces this difference.

Similarly, we propose that the number of rainy days is given by

$$D(m) = \frac{D_{max} - D_{min}}{2} \cos\left(\frac{2\pi}{12}(m - m_0)\right) + \frac{D_{max} + D_{min}}{2}, \quad (9)$$

where  $D_{max}$  and  $D_{min}$  correspond to the number of rainy days in the months labeled by  $m_0$  and  $m_0 + 6$ , respectively. Choosing  $m_0 = 2$ , this distribution would be suitable for the city of Córdoba. In Fig. 1(a) and (b) we show a schematic of the parameters used in Eqs. (8) and (9).

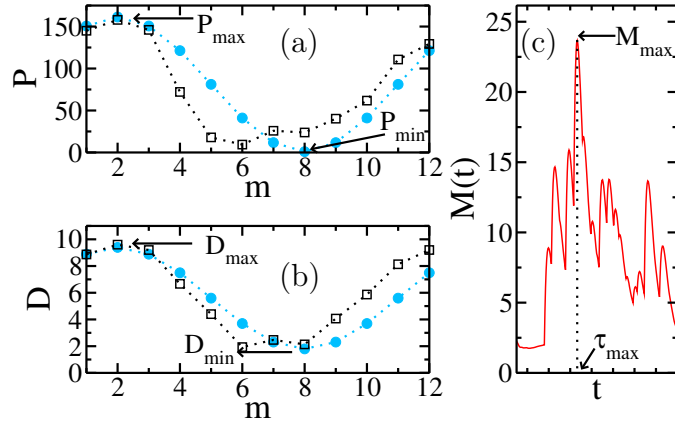


Figure 1: Schematic representation of Eq. (8) (panel (a)) and Eq. (9) (panel (b)), where  $P(m)$  is the monthly amount of rainfall and  $D(m)$  is the number of rainy days per month (blue circles). In these plots we also show the average monthly precipitation and number of rainy days in the city of Córdoba in the period 2001-2015 (black squares) (Wunderground (2016)). Panel (c) is a cartoon of  $M(t)$  indicating the maximum abundance of mosquitoes  $M_{max}$  and the time  $\tau_{max}$  (in day units) at which this peak is reached. Dotted lines are to guide the eye.

From Eqs. (8) and (9) we construct the amount of daily rainfall  $R(t)$  (see Eq. (4)),

placing the rainy days in each month at random and assuming that the amount of rain specified in Eq. (8) is equally distributed over these days. Then we calculate the factor  $\lambda(t)$  and integrate Eqs. (1) and (2). For simplicity we use in these equations the values of humidity and temperature obtained from meteorological data.

It is important to note that, by construction,  $R(t)$  is a stochastic time series since the rainy days for each month are chosen at random; therefore our results for the effect of the synthetic series  $R(t)$  on the mosquito abundance must be averaged over a large number (we take  $10^4$ ) of realizations. Using this model of time series of rainfall, we measure the highest peak of mosquito abundance  $M_{max}$  and the time  $\tau_{max}$  at which this peak is reached (see Fig. 1(c)).

In the following section, we explain how to introduce variability on the amount of daily rainfalls.

### *2.3. Model of synthetic rainfalls with variable daily intensity of precipitation*

In general, the daily rainfall intensity can range from drizzles with less than 1 mm to torrential downpours exceeding 200 mm (Li et al. (2013); Lei et al. (2008)). Various distributions, such as Weibull (Suhaila and Jemain (2007)), lognormal (Cho et al. (2004)) and generalized Pareto (Deidda (2010)), have been proposed to model the precipitation amount. While the above-mentioned distributions could be used to generate a sequence of rainfalls with variable or heterogeneous intensity, the disadvantage of this approach is that the total monthly rainfall is also a stochastic variable. In order to isolate the effect of the heterogeneity of the rainfalls, we will use a fracturing or fragmentation process (FT), which allows us to maintain the total monthly rainfall constant. This method is related to a cascading procedure that was used by physicists to study the fragmentation of brittle material (Hernandez (2003)). Recently the FT process was also applied to obtain empirical distributions with a finite tail (Finley and Kilkki (2014)). From a geometrical point of view (Borgos (2000); Finley and Kilkki (2014)), this process performs a sequential breakage of a segment or interval of length  $\ell$  to obtain  $D$  subintervals with variable length  $\tilde{\ell}$ , which can be used to decompose the total monthly rainfall into daily rainfalls with heterogeneous intensity. In this representation,  $\ell$  and  $\tilde{\ell}$  stand for the total amount of rainfall in a month and in a day, respectively. In Appendix B we explain the steps of the

FT process in detail.

This method depends on a parameter  $\alpha \in [0, 1]$  which controls the heterogeneity of the segment length. In particular:

- $\alpha = 0$  corresponds to the case where an interval is split into two subintervals of equal length  $\tilde{\ell} = \ell/2$ ,
- $\alpha = 1$  corresponds to the special case where two "intervals" are generated, one of length zero and the other of length  $\tilde{\ell} = \ell$ .

In Fig. 2 we show how  $\alpha$  controls the shape of the fragment length distribution  $\mathcal{P}(\tilde{\ell})$ , using  $\ell = 150$  mm and  $D = 10$ . Note that the resulting length of each subinterval corresponds to the intensity of rainfall in one day.

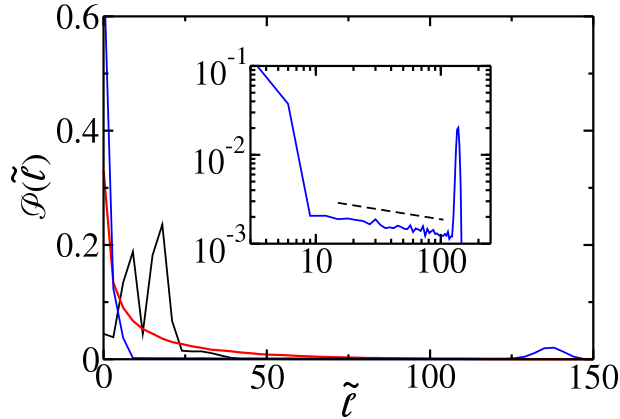


Figure 2: Distribution of length segments  $\mathcal{P}(\tilde{\ell})$ , using  $\ell = 150$  mm (total length) and  $D = 10$  (number of rainy days) for different values of  $\alpha$ : 0.1 (black), 0.5 (red), and 0.9 (blue). These distributions were obtained over  $2 \cdot 10^4$  realizations. In the inset, we show  $\mathcal{P}(\tilde{\ell})$  for  $\alpha = 0.9$  in log-log scale, and the dashed line corresponds to a power-law fit with an exponent 0.22.

For small values of  $\alpha$  the distribution is concentrated around the mean value close to  $\ell/D$ , while for intermediate values of  $\alpha$ , the distribution of lengths has a longer tail. Finally, for high values of  $\alpha$ ,  $\mathcal{P}(\tilde{\ell})$  has a peak near  $\ell$  which depicts a regime where most of the corresponding month total rainfall is confined to one day. Interestingly, we also note that in this case the rainfall distribution has a region in which it decays as a power law.

In order to model the temporal variation of precipitation  $R(t)$  (see Sec. 2.1.1), we apply a fracturing process (FT) for each month, partitioning an interval whose length is the amount of monthly precipitation given by Eq. (8) and where the number of subintervals (the number of rainy days) is given by Eq. (9). See Appendix B for further details on the construction of  $R(t)$ .

### 3. Results

#### 3.1. Calibration

We calibrate our model of mosquito abundance using a Metropolis-Hastings algorithm (see Appendix A) with the number of female *Culex quinquefasciatus* mosquitoes collected in Córdoba city (31°24'30" S, 64°11'02" W, Córdoba province, Argentina) every two weeks from January 2008 to December 2009 (see Batallán (2013); Batallán et al. (2015) for details on the data and their sources). The climate of Córdoba is temperate with dry winters and hot rainy summers. The mean annual temperature ranges between 16°C-17°C and the mean annual rainfall is 800 mm (Jarsún et al. (2003)). The temperature  $[T(t)]$ , relative humidity  $[Hum(t)]$ , and rainfall  $[R(t)]$  data for Córdoba were obtained from the website Wunderground (2016). We calibrate the following parameters:  $\beta_L$ ,  $H_{max}$ ,  $H_{min}$ , and  $K_L$ . In Appendix C we perform a sensitivity analysis of these calibrated parameters.

As initial conditions of Eqs. (1) and (2), we set  $M(t) = L(t) = 20$ . To attenuate the effect of these initial conditions, the integration of the equations starts 12 months before we implement the fitting and study our model.

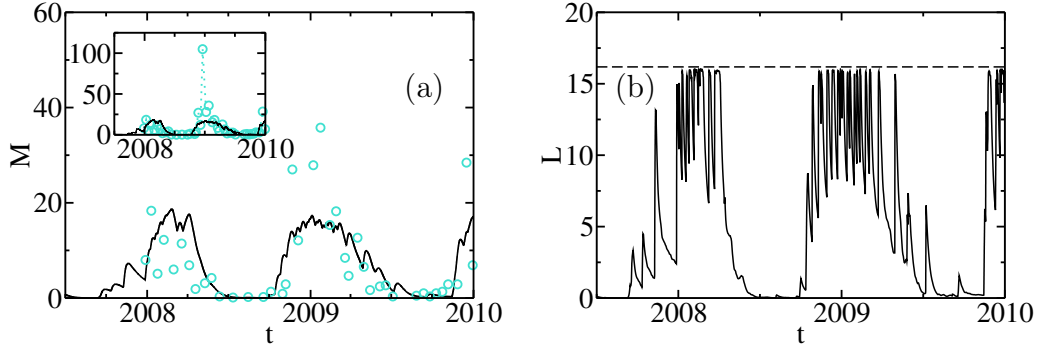


Figure 3: Panel (a): Evolution of the abundance  $M(t)$  of adult mosquitoes obtained from the fitted compartmental model. Large tick marks correspond to Jan 1. The blue circles represent the number of female mosquitoes collected in the city of Córdoba, and the black line corresponds to the model fit. Inset: main plot on a larger vertical scale showing an extremely high abundance data point. Panel (b): Evolution of the number of immature mosquitoes obtained from Eqs. (1)-(7). The dashed line represents the carrying capacity,  $K_L$ .

Fig. 3(a) shows the fit of our model to the data, where the abundance  $M(t)$  is given as the number of female mosquitoes per night per trap. We observe that the mosquito population, which is assumed to be proportional to  $M(t)$ , increases in summer as expected. Although there are no daily abundance data to compare with, we remark that our model predicts day-to-day changes in the abundance of adult mosquitoes  $M(t)$  due to fluctuations in temperature, humidity, and rainfall.

In Fig. 3(b) we plot the evolution of the immature population of mosquitoes  $L(t)$  which shows that temperature and precipitation lead to more abrupt fluctuations in this group than in the compartment of adult mosquitoes. This was to be expected, since these climatic variables are directly introduced into the equation of the population of immature mosquitoes (see Eq. (1)). In turn, we find that after a rainfall event there is a large increase in the abundance of this group which frequently approaches the carrying capacity, leading to a subsequent population decline due to competition among immature mosquitoes (Roberts and Kokkinn (2010); Suleman et al. (1982)).

In the following section we study how different rainfall patterns affect the mosquito population dynamics.

### 3.2. Effect of different rainfall regimes on $M_{max}$

The proposed model of synthetic rainfall allows us to explore the evolution of mosquito abundance under possible scenarios in which the weather becomes, for instance, rainier, or with persistent drought conditions. In this Section we discuss how different rain regimes influence the mosquito population when it is at its highest (see Fig. 1(c)). To do this, we first assume that the total rainfall  $P(m)$  (see Eq. (8)) is equally distributed in  $D(m)$  days (see Eq. (9)), the remainder of the days in the month being rainless.

In Fig. 4(a) we consider the weather data (temperature and humidity) of the austral summer season 2008-2009, to show the influence of  $P_{max}$  and  $D_{max}$  on the highest peak  $M_{max}$  of mosquito abundance, with fixed values of the parameters  $P_{min} = 10$  mm and  $D_{min} = 1$ .

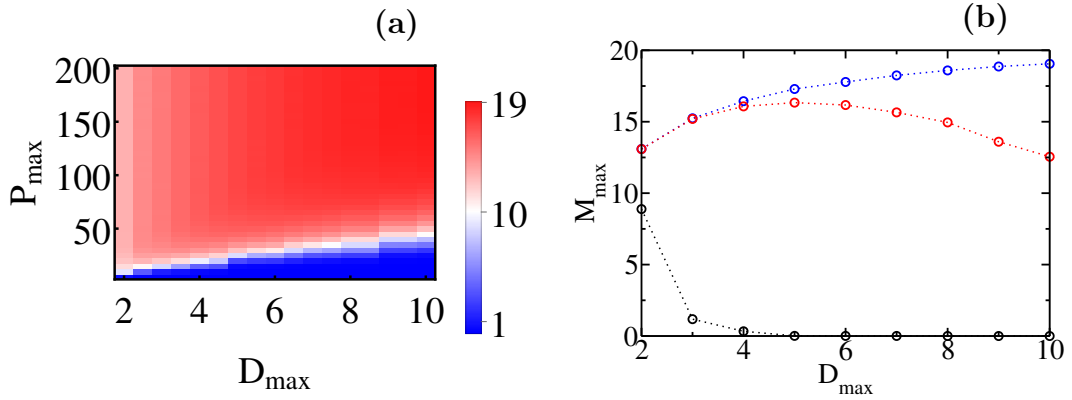


Figure 4: Maximum number of mosquitoes  $M_{max}$  in the plane  $P_{max}$ - $D_{max}$  (a) and  $M_{max}$  as a function of  $D_{max}$  (b) for  $P_{max} = 10$  mm (black),  $P_{max} = 50$  mm (red) and  $P_{max} = 190$  mm (blue). Dotted lines are to guide the eye. The results were obtained from Eqs. (1)-(7) and  $10^4$  stochastic realizations.

From Fig. 4(a), we note that, for a fixed number of rainy days  $D_{max}$ , the highest peak of mosquito abundance increases as  $P_{max}$  grows since, as expected, a greater amount of water promotes breeding sites for mosquitoes. Similarly, it can be seen from Figs. 4(a) and (b) that for  $P_{max} \approx 190$  mm,  $M_{max}$  is an increasing function with  $D_{max}$ , because a higher frequency of rainfall events provides more opportunities for mosquitoes to breed. The opposite happens for the lowest level of precipitation ( $P_{max} \approx 10$  mm) as there is very little daily rainfall in this regime and the accumulated water evaporates quickly. Interestingly, we find that at moderate precipitation levels  $M_{max}$  has a maximum at an intermediate value of the number of rainy days. If the rainfall is equally distributed over a

few days, the mosquito population will increase with more rainy days, but, beyond certain point, the rain becomes too thin to maintain all breeding sites active and the mosquito population must decrease. However, as it was mentioned above, the intensity of daily rainfalls is usually far from uniform (Wunderground (2016)), and recent studies suggest that the amount of mosquitoes depend on the distribution of precipitation (Cheng et al. (2016); Wang et al. (2016); Bomblies (2012); Bomblies et al. (2008)). Therefore, in the following we study how heterogeneity in the daily rainfall affects the dynamics of mosquito abundance.

In figure 5, using the weather data of the austral summer season 2008-2009 (temperature and humidity), we show the peak mosquito abundance for different values of  $P_{max}$ ,  $D_{max}$ , and  $\alpha$ , as obtained from the Eqs. (1)-(7). For simplicity, we use the same value of  $\alpha$  for each month.

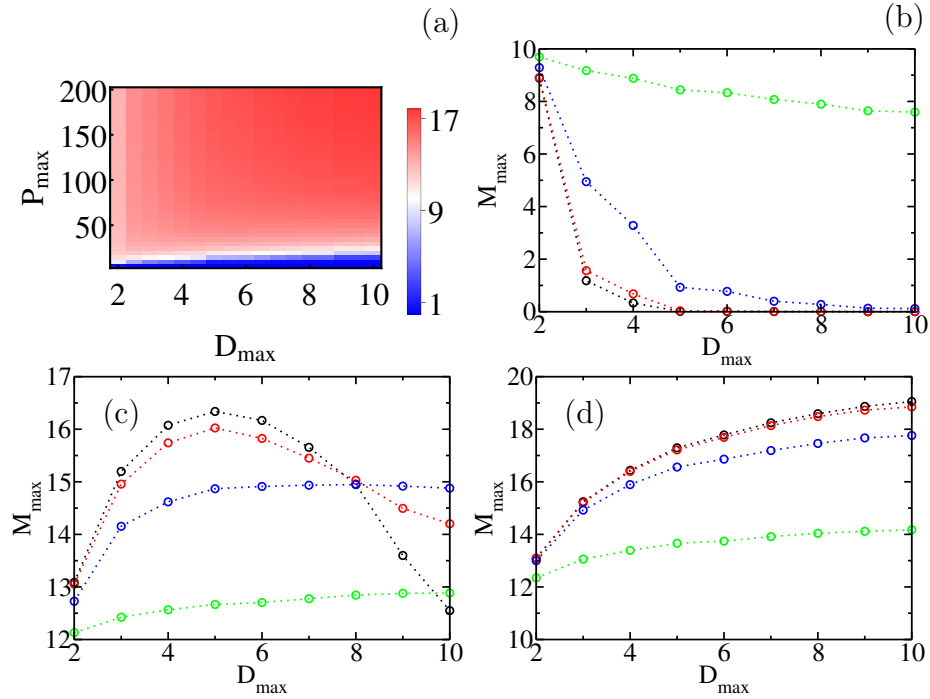


Figure 5: Panel (a): Maximum number of mosquitoes  $M_{max}$  in the plane  $P_{max}$ - $D_{max}$  for  $\alpha = 0.5$  and fixed values of the parameters  $D_{min} = 1$  and  $P_{min} = 10$  mm. Panel (b):  $M_{max}$  as a function of the number  $D_{max}$  of rainy days in the wettest month for  $P_{max} = 10$  mm for different values of  $\alpha$ . Rainy days have either the same amount of rainfall intensity (black), or  $\alpha = 0.1$  (red),  $\alpha = 0.5$  (blue), or  $\alpha = 0.9$  (green). Panels (c) and (d) show the same as (b) for  $P_{max} = 50$  mm and  $P_{max} = 190$  mm, respectively. Averaging was made over  $10^4$  realizations of the rainfall time series  $R(t)$ .

From Figs. 5(a), (b) and (d), we note that, similarly to the case of constant rainfall intensity, the abundance  $M_{max}$  is a decreasing (increasing) function with  $D_{max}$  for a low (high) amount of monthly precipitation  $P_{max}$ . Furthermore, Figs.5(b)-(d) show that a higher variability in the intensity of rainfalls reduces the dependence of  $M_{max}$  with  $D_{max}$  with respect to the case of homogeneous rainfall. In particular, for  $P_{max} = 10$  mm (see Fig.5(b)), we note that  $M_{max}$  is higher in the heterogeneous case ( $\alpha > 0$ ) than in the homogeneous one, since a greater variability tends to confine most of the total monthly precipitation to a few days, in which there is a higher level of water (see Eq. (6)) available for the immature mosquito population. In contrast, for  $P_{max} = 190$  mm (see Fig.5(d)) the heterogeneity diminishes the abundance  $M_{max}$  with respect to the homogeneous one. In this case, even if a higher amount of precipitation in a few days would increase the rate of immature mosquito birth, the effect of the intense rain days is limited by the threshold  $H_{max}$  because the impact of precipitation saturates for a precipitation higher than  $H_{max}$  (*i.e.*,  $\lambda(t) = 1$ , see Eqs. (6) and (7)). Moreover, there are more rainy days with low precipitation in this regime which further reduces the growth of the mosquito population. As a consequence, the heterogeneity in the intensity of rainfall implies that the monthly total precipitation is a more relevant variable for the prediction of the maximum abundance of mosquitoes than the number of rainy days.

Another aspect of relevance for the prediction of vector-borne diseases is the effect of the rainfall in the dry-season (winter) on the future abundance of mosquitoes in summer. To study this relationship, we measure the maximum abundance of mosquitoes  $M_{max}$  and the timing  $\tau_{max}$  at which the peak of mosquito abundance is reached (see Fig. 1(c)), for different values of  $P_{min}$  and  $D_{min}$ , which correspond to the parameters that control the intensity and frequency of rainfall in the driest month, respectively. Here, we keep fixed the parameters  $P_{max} = 150$  mm and  $D_{max} = 10$ , and assume that February is always the wettest month of the year ( $m_0 = 2$ , see Eq. (8)).



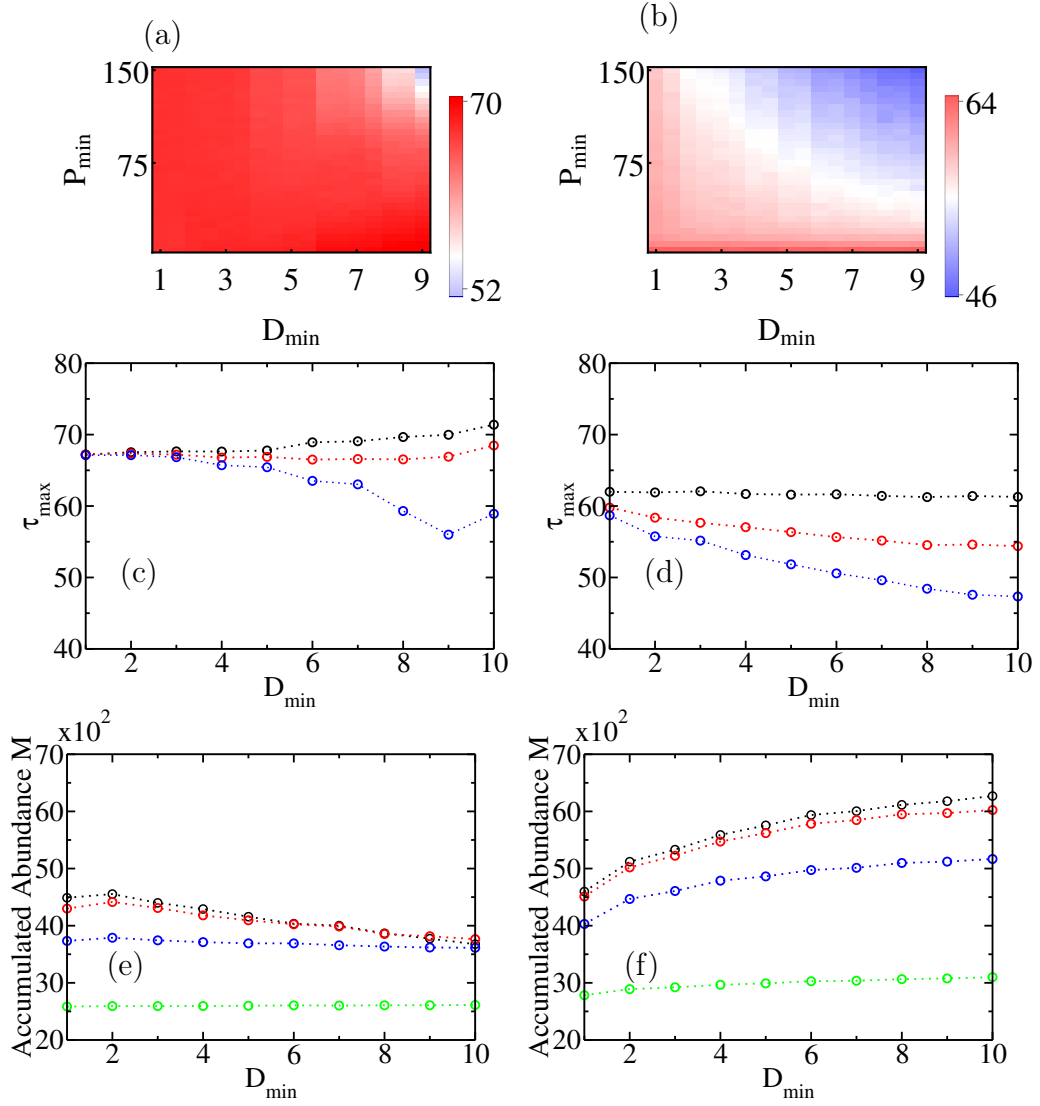


Figure 6: Panels (a) and (b): Time  $\tau_{max}$  (measured in days, where  $\tau_{max} = 0$  corresponds to January 1, 2009) at which the peak of mosquito abundance is reached in the plane  $P_{min}$ - $D_{min}$  for a homogeneous intensity distribution of rainfalls (a) and  $\alpha = 0.9$  (b). Panels (c) and (d):  $\tau_{max}$  as a function of the number of rainy days in the driest month,  $D_{min}$ , for a homogeneous intensity distribution of rainfalls (c) and  $\alpha = 0.9$  (d) for various values of  $P_{min}$ : 10 mm (black), 50 mm (red) and 130 mm (blue). Panels (e) and (f): accumulated mosquito abundance between July 2008 and July 2009 as a function of  $D_{min}$ , with  $P_{min} = 10$  mm (e) and  $P_{min} = 130$  mm (f) for: homogeneous rainfall intensity (black),  $\alpha = 0.1$  (red),  $\alpha = 0.5$  (blue) and  $\alpha = 0.9$  (green). The dotted lines are a guide to the eye. The figures were obtained averaging over  $10^5$  realizations of the rainfall time series  $R(t)$ .

We note from Figs. 6(a) and (c) that, in the case of a homogeneous distribution of rainfalls, the time of peak  $\tau_{max}$  moves forward for high values of  $P_{min}$  and  $D_{min}$ , because in this case the rainfalls are regular and abundant throughout the year, which favors

mosquito breeding. However, for the explored values of  $P_{min}$  and  $D_{min}$ , the position of this peak is in late February or March, *i.e.*, just after the wettest month of the year. On the other hand, in a scenario of heterogeneous rainfalls with  $\alpha = 0.9$  (see Figs. 6(b) and (d)),  $\tau_{max}$  is moved forward by only approximately 10 days with respect to the homogeneous intensity case. Correspondingly, for constant values of  $\alpha$  (0.1, 0.5 and 0.9) the changes in  $P_{min}$  and  $D_{min}$  only affect  $M_{max}$  by less than 5% (not shown here). Therefore these results suggest that the peak of abundance of mosquitoes and  $\tau_{max}$  are mainly determined by summer weather conditions and the carrying capacity of the system and not by the intensity and distribution of precipitation throughout the year. Despite the weak effect of  $P_{min}$  on  $M_{max}$ , Figs. 6 (e) and (f) show, as expected, that an increasing value of  $P_{min}$  could have a remarkable effect on the accumulated abundance of mosquitoes (one measure of which is the time integral of  $M(t)$  over the period of interest), since from  $P_{min} = 10$  mm to  $P_{min} = 130$  mm, it could increase by more than 40%. However, for the case of a fixed value of  $P_{min}$  and higher values of  $\alpha$  we obtain that the accumulated abundance of mosquitoes diminishes down to a 50% of the value for a homogeneous rain distribution. Consequently, the heterogeneity could help to attenuate the enhancement of the mosquito population. Therefore, these findings suggest that in order to predict the total annual abundance, it is not only necessary to take into account the overall amount of rainfall throughout the year but also the heterogeneity in daily rainfall intensity.

#### 4. Discussion

Since projections of climate change (Nuñez et al. (2009)) suggest that for the late twenty-first century in regions of South America, such as Córdoba province, the pattern of precipitation will change towards a regime with rainier autumns and an increase in extreme events, it is crucial to study how this variation would affect the mosquito abundance.

In this paper we studied the effects of the total intensity, number of rainy days and heterogeneity of rainfall on the mosquito population. We found that for a regime with a low total rainfall, the abundance of mosquitoes is a decreasing function with the number of rainy days, while for a high total rainfall regime it is an increasing function of this number. Interestingly, for an intermediate precipitation regime, we found that there is a halfway number  $D_{max}$  of rainy days for which  $M_{max}$  is optimized. Since  $P_{max}$  is

fixed, fewer rainy days would imply dry intervals, leading to a lessening of the mosquito abundance. If the number of rainy days exceeds the optimal value of  $D_{max}$ , a considerable fraction of the rainwater resulting from the typically meager rainfall would disappear due to evapotranspiration, again leading to a reduction of the mosquito abundance.

In order to study the effect of the heterogeneity in the daily rainfall, we used a fracturing process that keeps constant the total amount of monthly precipitation. We observed that a higher heterogeneity reduces the dependence of  $M_{max}$  on the number of rainy days. However, an increasing variability favors the mosquito production in the low rainfall regime, while the opposite behavior takes place in the case of high precipitation  $P_{max}$ . Therefore, if climatic models predict the intensification of storms, but not an increase in the total amount of precipitation, our model predicts that the enhancement of mosquito abundance would be more significant in semiarid areas than in humid climates.

Finally we study the effect of an increasing amount of rainfall in the dry season on the mosquito abundance dynamics, obtaining that high precipitation throughout the year does not significantly alter the maximum abundance or the time at which this peak occurs, but it could notably increase the accumulated abundance of mosquitoes. However, we also observed that a regime with a higher variability of rainfall intensity could reduce this increase.

While our model captures multiple relationships between rainfall and mosquito population, additional extensions could be considered. For instance, there is evidence that rainfalls reduce the immature population in the short term due to flushing of breeding sites (Gardner et al. (2012); Strickman (1988)) and affect the bacterial concentration used as food by mosquito larvae (Chaves and Kitron (2011)); therefore, it would be interesting to study the relevance of these effects on the dynamic of mosquito population.

We think that our findings could be used as support and reference guidance for the assessment of the influence of different rainfall regimes on the mosquito population dynamics, using the weather data for any specific region. Such an assessment would impact positively on our ability to make predictions for the spread of various possible arboviruses. It is also known that rainfall could have a substantial effect on insecticide residence times (Allan et al. (2009)). The model presented here can be used to optimize

the efficacy of mosquito control campaigns, using temperature and rainfall data to select the best times for the application of population reduction procedures.

## Appendix A. Calibration

The Metropolis-Hastings (MH) algorithm is a stochastic optimization tool for fitting statistical models to data that has been used in cosmology (Christensen et al. (2001, 2004); Lewis and Bridle (2002)), epidemiology (Merler et al. (2015)), and in the study of mosquito population dynamics (Marini et al. (2016)). This algorithm allows us to estimate the unknown values of some parameters  $\Theta$  ( $\Theta$  represents either a single parameter or a parameter set), by means of a stochastic search in the parameter space that generates a sequence or chain  $\Theta^{(i)}$ , where  $i$  represents the step number of the MH algorithm (Bonamente (2013)). Each value of this chain is sampled from a proposal distribution and accepted with a probability  $\sigma$  defined by an acceptance function, which depends on the likelihood function of the observations.

In our model we estimate the parameters  $\beta_L$ ,  $H_{max}$ ,  $H_{min}$  and  $K_L$  using a MH algorithm and the data for adult mosquito abundance from Córdoba city in the period 2008-2009 (Batallán (2013); Batallán et al. (2015)). We propose that the likelihood of the observations is given by

$$L = \prod_{j=1}^n p(x_j(H_{max}, H_{min}, K_L, \beta_L); k_j), \quad (\text{A.1})$$

where  $n$  is the number of data points and  $p(x_j(H_{max}, H_{min}, K_L, \beta_L); k_j)$  is the probability to observe the abundance  $k_j$  of mosquitoes obtained from the data. Here we assume that  $p(\cdot)$  follows a Poisson distribution whose mean  $x(H_{max}, H_{min}, K_L, \beta_L)$  is the number of adult mosquitoes predicted by Eqs. (1)-(7).

The MH algorithm implemented in this paper has the following steps:

- Step 1: Initialize the starting value of the parameters  $\Theta^{(i=0)}$ , using a uniform distribution in order to avoid favoring any initial value.
- Step 2: Generate a new sample of the parameters,  $\Theta^{New}$  starting from a proposal distribution that indicates a candidate for the next sample value. To ensure that

the new values of the parameters are positive, we use as a proposal distribution a log-normal density which has a mean equal to the logarithm of the current value parameter and constant variance  $\delta$ . The value of this variance is chosen in order to guarantee an acceptance rate between 10% and 30% in the burn-in period.

- Step 3: accept the new candidate  $\Theta^{New}$  with probability  $\sigma$ :

$$\sigma = \min \left\{ 1, \frac{L(\Theta^{New})}{L(\Theta^{(i)})} \right\}. \quad (\text{A.2})$$

- Step 4: repeat steps 2 and 3 until convergence is reached.

We perform  $2 \cdot 10^6$  iterations and check convergence by visual inspection of the chain  $\Theta^{(i)}$ . In order to construct the posterior distribution of the parameters, we discard the first  $10^5$  iterations as a burn-in and we only keep every  $20^{th}$  sampled value of the remaining iterations to reduce autocorrelation within successive samples. Finally, the values of the parameters that we will use in our model are the averages of the medians of the posterior distributions obtained from 5 different initial conditions (see step one of the MH algorithm).

Fig. A.1 shows the posterior distribution obtained for the parameters  $\beta_L$ ,  $H_{max}$ ,  $H_{min}$  and  $K_L$ . We note that all of these distributions are unimodal, except for  $H_{max}$ . Although in this paper we set  $H_{max} = 9.86$  mm, since it is the average value of the median obtained from the MH algorithm, we also check our model for  $H_{max} \approx 7$  mm and  $H_{max} \approx 13$  mm which are the positions of the highest peaks of the posterior distribution (see Fig. A.1(b)). For these cases, our results presented in Section 3 do not qualitatively change.

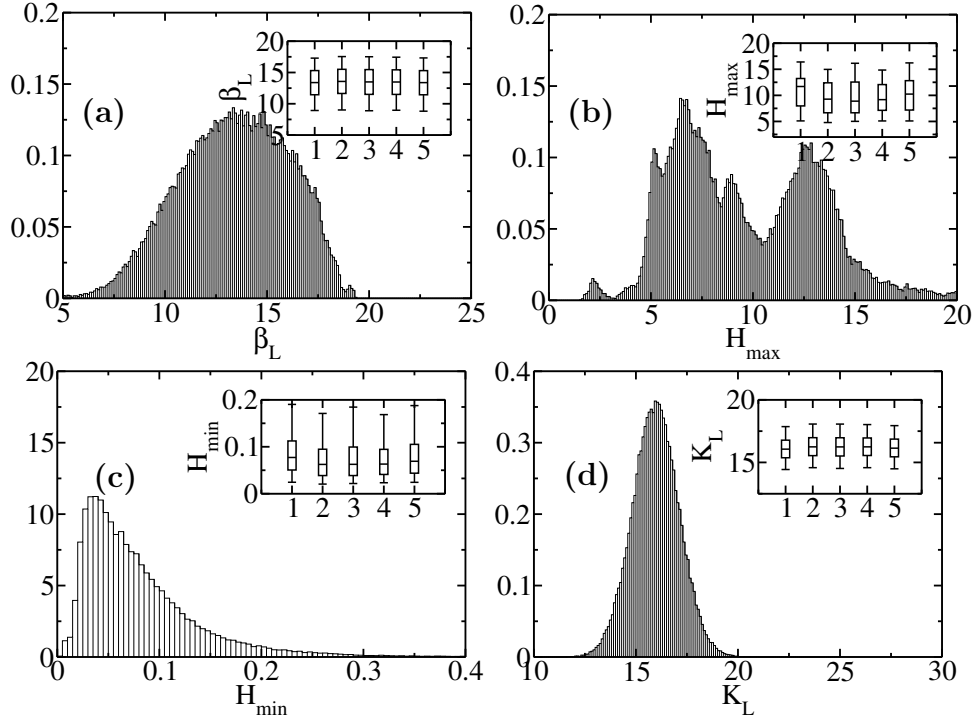


Figure A.1: Posterior distribution obtained in the steady state of the MH algorithm from  $2 \cdot 10^6$  iterations, using  $10^5$  iterations as a burn in for the parameters  $\beta_L$  (a),  $H_{max}$  (b),  $H_{min}$  (c), and  $K_L$  (d). In the insets we show the boxplot obtained from the posterior distribution in the steady state for 5 different initial conditions (see step one of the MH algorithm).

## Appendix B. Fracturing process

Fracturing process (FT) is a stochastic iterative process which generates a finite sequence of numbers with the property that their sum is always a constant (Borgos (2000); Finley and Kilkki (2014)). From a geometrical point of view, this method consists of partitioning an interval of length  $\ell$  in a number  $D$  of subintervals or segments, with the property that the sum of their lengths is always  $\ell$ . Following Finley and Kilkki (2014), the FT process starts with an interval or segment of length  $\ell$  which is split into two subintervals of lengths  $\tilde{\ell}$  and  $\ell - \tilde{\ell}$ , where  $\tilde{\ell}$  is a stochastic variable generated by the following function:

$$\tilde{\ell} = \ell \times \begin{cases} \rho^{\frac{1-\alpha}{\alpha}} & \text{if } 0 \leq \rho < \frac{\alpha}{2} \\ \frac{1}{2} + \left(\rho - \frac{1}{2}\right) \frac{\alpha}{1-\alpha} & \text{if } \frac{\alpha}{2} \leq \rho \leq 1 - \frac{\alpha}{2} \\ 1 - (1 - \rho)^{\frac{1-\alpha}{\alpha}} & \text{if } 1 - \frac{\alpha}{2} < \rho \leq 1 \end{cases} \quad (\text{B.1})$$

Here  $\rho$  is a uniform random variable and  $\alpha \in (0, 1)$  is a parameter that controls the average length  $\tilde{\ell}$ . In a second step, the partition function, Eq. (B.1), is applied again on the intervals resulting from the previous step, generating a total of 4 intervals. This procedure is repeated until the required number of intervals is reached <sup>1</sup>.

In order to model the variability in the daily rainfall  $R(t)$ , we apply a FT process for each month, in which,

- the length of the initial segment  $\ell$  is given by Eq. (8), *i.e.*, the monthly precipitation,
- the number  $D$  of subintervals is given by Eq. (9), *i.e.*, the number of rainy days.

After we apply the fracturing process, the length of each resulting interval  $\tilde{\ell}_i$  (with  $i = 1, \dots, D$ ) represents the total amount of water that falls in the day  $d_i$ , which we choose at random as it is shown in the schematic of Fig. B.1, and then we set  $R(d_i) = \tilde{\ell}_i$ .

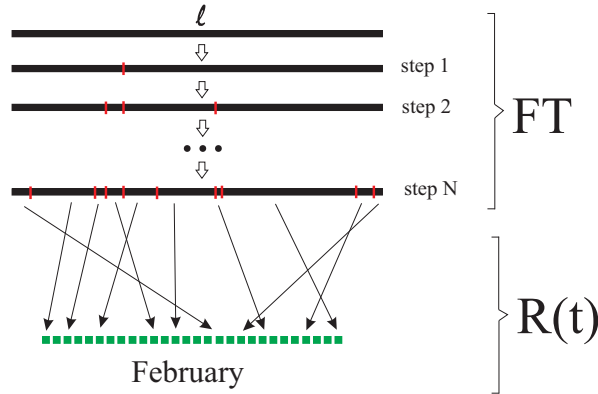


Figure B.1: Schematic of the construction of  $R(t)$  corresponding to February. At the top of the figure we show the FT process and at the bottom the construction of  $R(t)$ , in which each square represents a day in February. The FT process stops when the required number of subintervals (given by Eq. (9)) is reached. The length of each of these segments represents the rainfall in one day in February, which is randomly chosen. Those days that are not associated with any length of the FT process, have  $R(t) = 0$ .

In Fig. B.2 we plot the distribution of segment lengths obtained from an FT process

<sup>1</sup>For example, in order to generate a total of five subintervals, three iterations of the fracturing process must be performed: step 1) the initial interval is divided into two subintervals, step 2) each of the previous subintervals is divided into two parts, and finally step 3) one randomly chosen subinterval of the previous step is split into two subintervals.

for  $\ell = 150$  mm and  $D = 10$ , which are the average rainfall and the number of rainy days in February, respectively. Here we use the value of  $\alpha = 0.43$  which gives the best fit to the February rainfalls in the city of Córdoba in the period 2001-2015.

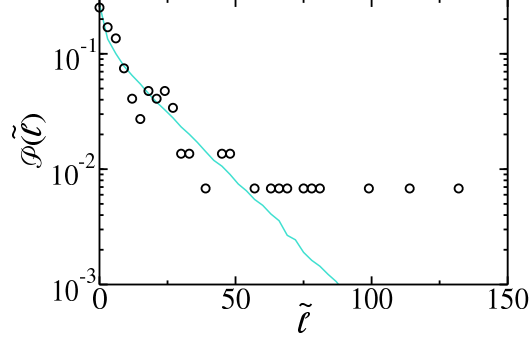


Figure B.2: Observed February rainfall distribution in Córdoba during the period 2001-2015 (symbols) and distribution of segment lengths  $\mathcal{P}(\tilde{\ell})$  for  $\ell = 150$  mm, and  $D = 10$ , and  $\alpha = 0.43$  (solid line) obtained over  $2 \cdot 10^4$  realizations.

## Appendix C. Sensitivity analysis

A sensitivity analysis allows us to measure the impact of different parameters on the relevant variables of our model. In this section, we perform a one-way sensitivity analysis on the model of Sec. 2.1, by varying a  $\pm 25\%$  of the baseline values of individual parameters one at a time, while keeping the other parameters constant in order to analyze their individual impact on the maximum abundance of mosquitoes  $M_{max}$ . The parameters examined are:  $\beta_L$ ,  $H_{max}$ ,  $H_{min}$ , and  $K_L$ . The results of the sensitivity analysis are summarized in Table C.1.

Table C.1: Variation of the maximum abundance of mosquitoes ( $M_{max} = 17.3$ ) when it is applied a one-way sensitivity analysis.

Parameter	-25%	+25%
$\beta_L$	-2.1%	+1%
$H_{max}$	-0.4%	+0.9%
$H_{min}$	-0.7%	+0.6%
$K_L$	-25%	+25%



It shows, as expected, that for higher values of  $\beta_L$ ,  $H_{max}$ ,  $H_{min}$ , and  $K_L$  the abundance  $M_{max}$  increases. Although the influence of the first three is rather weak,  $M_{max}$  is heavily influenced by  $K_L$ , which is therefore a critical parameter for the estimation of mosquito abundance.

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